

AMENDMENTS TO THE CLAIMS

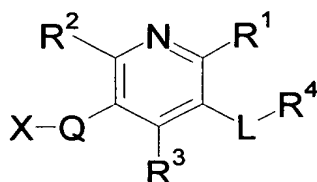
1. (Cancelled)
2. (Cancelled)
3. (withdrawn) The compound of claim 1, wherein the acyl group for X is a carboxyl group.
4. (Cancelled)
5. (Cancelled)
6. (Withdrawn) The compound of claim 1, wherein R⁴ is an amino group.
7. (Withdrawn) The compound of claim 1, wherein L is a C₁₋₁₀ alkylene group.
8. (Cancelled)
9. (Cancelled)
10. (Cancelled)
11. (Cancelled)
12. (Withdrawn) A prodrug of a compound of claim 1 or a salt thereof.
13. (Cancelled)
14. (Cancelled)
15. (Cancelled)
16. (Cancelled)
17. (Cancelled)
18. (Cancelled)

19. (Cancelled)

20. (Cancelled)

21. (Cancelled)

22. (New) A compound represented by the formula



wherein

R¹ and R² are the same or different and each is

(1) a C₁₋₁₀ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a C₃₋₁₀ cycloalkyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkoxy group;

(2) a C₆₋₁₄ aryl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

(3) a C₇₋₁₃ aralkyl group;

R³ is a C₆₋₁₄ aryl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s), a halogen atom, a C₁₋₆ alkoxy-carbonyl group, a carboxyl group, a hydroxy group, and a C₁₋₆ alkoxy group optionally substituted by 1 to 3 halogen atom(s);

R⁴ is an amino group;

L is a C₁₋₁₀ alkylene group;

Q is a bond, a C₁₋₁₀ alkylene group or a C₂₋₁₀ alkenylene group; and

X is

(2) a cyano group;

(3) (3a) a carboxyl group;

(3b) a carbamoyl group;

(3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3

substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy group;

_____ (3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3f) a C₇₋₁₃ aralkyloxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group, a halogen atom, a cyano group, a nitro group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom and a C₁₋₆ alkoxy group;

_____ (3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

_____ (3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3m) a C₁₋₆ alkylsulfonyl group optionally substituted by 1 to 3

substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3n) a C₆₋₁₄ arylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkylsulfonyl group;

_____ (3o) a nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3p) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3q) a C₇₋₁₃ aralkyl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3r) a non-aromatic heterocycloxy-carbonyl group;

_____ (3s) a phosphono group optionally mono- or di-substituted by a C₁₋₆ alkyl group;

_____ (3t) an aromatic heterocyclyl-C₇₋₁₃ aralkyloxy-carbonyl group;

_____ (3u) a C₃₋₁₀ cycloalkyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (3v) a C₆₋₁₄ aryl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from an amino group optionally mono- or di-substituted by a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group; or

_____ (3w) an aromatic heterocyclyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4) (4a) a C₁₋₆ alkyl-carbonyloxy group;

_____ (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (4d) a 5- or 6-membered aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4e) a fused aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4f) an aromatic heterocyclyl-C₁₋₆ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (4g) an aromatic heterocyclyl-C₆₋₁₄ aryloxy group;

_____ (5) (5a) a C₁₋₆ alkylthio group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (5b) a C₆₋₁₄ arylthio group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group and a carbamoyl group; or

_____ (5c) a 5- or 6-membered aromatic heterocyclylthio group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6) (6a) an amino group;

_____ (6b) a C₁₋₆ alkoxy-carbonyl-C₁₋₁₀ alkylamino group;

_____ (6c) a carboxy-C₁₋₁₀ alkylamino group;

_____ (6d) a C₇₋₁₃ aralkyloxy-carbonylamino group optionally substituted

by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6e) a carbamoylamino group;

_____ (6f) a mono- or di-C₁₋₆ alkyl-carbamoylamino group;

_____ (6g) a C₁₋₆ alkylsulfonylamino group;

_____ (6h) a C₆₋₁₄ arylsulfonylamino group optionally substituted by a C₁₋₆ alkylsulfonyl group;

_____ (6i) an aromatic heterocyclyl-sulfonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group and a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

_____ (6j) a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

_____ (6k) a C₃₋₁₀ cycloalkyl-carbonylamino group;

_____ (6l) a C₆₋₁₄ aryl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a cyano group, an optionally halogenated C₁₋₆ alkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group;

_____ (6m) a C₇₋₁₃ aralkyl-carbonylamino group;

_____ (6n) a C₈₋₁₃ arylalkenyl-carbonylamino group;

_____ (6o) an aromatic heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a C₆₋₁₄ aryl group, a C₇₋₁₃ aralkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6p) a nitrogen-containing heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6q) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonylamino group;

_____ (6r) a tetrahydropyranylcabonylamino group;

_____ (6s) a 4-oxo-4,5,6,7-tetrahydro-1-benzofuranyl-carbonylamino group;

_____ (6t) a C₁₋₆ alkoxy-carbonylamino group optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (6u) a C₆₋₁₄ aryloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6v) a C₇₋₁₃ aralkyl-carbamoylamino group; or

_____ (6w) an aromatic heterocyclyl-carbamoylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (7) (7a) tetrazolyl;

_____ (7b) oxoimidazolidinyl;

_____ (7c) dioxoimidazolidinyl optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7d) oxopiperazinyl;

_____ (7e) dioxopiperazinyl;

_____ (7f) oxodihydrooxadiazolyl;

_____ (7g) dioxoisindolyl;

_____ (7h) oxazolyl optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (7i) dioxooxazolidinyl or dioxothiazolidinyl, each of which is optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7j) 4-oxo-2-thioxo-1,3-thiazolidin-5-yl or 4-oxo-2-thioxo-1,3-oxazolidin-5-yl, each of which is optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7k) 1,3(2H,5H)-dioxo-tetrahydroimidazo[1,5-a]pyridinyl;

_____ (7l) 1,3(2H,5H)-dioxo-10,10a-dihydroimidazo[1,5-b]isoquinolinyl; or

_____ (7m) a C₆₋₁₄ aryl group optionally substituted by a C₁₋₆ alkoxy-

carbonyl group;

provided that

when X is an ethoxycarbonyl group, then Q is a C₁₋₁₀ alkylene group or a C₂₋₁₀ alkenylene group
or a salt thereof.

23. (New) The compound of claim 22, wherein X is

(2) a cyano group;

(3) (3a) a carboxyl group;

(3b) a carbamoyl group;

(3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3
substituent(s) selected from a carboxyl group, a carbamoyl group, a
thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy
group;

(3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally
substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl
group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group
optionally substituted by a C₁₋₆ alkyl group;

(3f) a C₇₋₁₃ aralkyloxy-carbonyl group optionally substituted by 1 to
3 substituent(s) selected from a carboxyl group, a carbamoyl group, a
thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group, a halogen atom, a cyano
group, a nitro group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group and a C₁₋₆
alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s)
selected from a halogen atom, a carboxyl group, C₁₋₆ alkoxy-carbonyl group and
a carbamoyl group);

(3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl
group optionally substituted by 1 to 3 substituent(s) selected from a halogen
atom and a C₁₋₆ alkoxy group;

(3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-
substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

(3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally

substituted by a C₁₋₆ alkyl group;

_____ (3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

_____ (3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3m) a C₁₋₆ alkylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3n) a C₆₋₁₄ arylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkylsulfonyl group;

_____ (3o) a nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3p) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3q) a C₇₋₁₃ aralkyl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3r) a non-aromatic heterocycloxy-carbonyl group;

_____ (3s) a phosphono group optionally mono- or di-substituted by a C₁₋₆ alkyl group;

_____ (3t) an aromatic heterocyclyl-C₇₋₁₃ aralkyloxy-carbonyl group;

_____ (3u) a C₃₋₁₀ cycloalkyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (3v) a C₆₋₁₄ aryl-carbamoyl group optionally substituted by 1 to 3

substituent(s) selected from an amino group optionally mono- or di-substituted by a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group; or

_____ (3w) an aromatic heterocyclyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4) (4a) a C₁₋₆ alkyl-carbonyloxy group;

_____ (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (4d) a 5- or 6-membered aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4e) a fused aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4f) an aromatic heterocyclyl-C₁₋₆ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (4g) an aromatic heterocyclyl-C₆₋₁₄ aryloxy group;

_____ (5) (5a) a C₁₋₆ alkylthio group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (5b) a C₆₋₁₄ arylthio group optionally substituted by 1 to 3
substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆
alkylthio group and a carbamoyl group; or

_____ (5c) a 5- or 6-membered aromatic heterocyclylthio group optionally
substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl
group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (7) (7a) tetrazolyl;

_____ (7b) oxoimidazolidinyl;

_____ (7c) dioxoimidazolidinyl optionally substituted by a C₁₋₆ alkyl group
optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and
a C₁₋₆ alkoxy-carbonyl group;

_____ (7d) oxopiperazinyl;

_____ (7e) dioxopiperazinyl;

_____ (7f) oxodihydrooxadiazolyl;

_____ (7g) dioxoisindolyl;

_____ (7h) oxazolyl optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (7i) dioxooxazolidinyl or dioxothiazolidinyl, each of which is
optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3
substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7j) 4-oxo-2-thioxo-1,3-thiazolidin-5-yl or 4-oxo-2-thioxo-1,3-
oxazolidin-5-yl, each of which is optionally substituted by a C₁₋₆ alkyl group
optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and
a C₁₋₆ alkoxy-carbonyl group;

_____ (7k) 1,3(2H,5H)-dioxo-tetrahydroimidazo[1,5-a]pyridinyl;

_____ (7l) 1,3(2H,5H)-dioxo-10,10a-dihydroimidazo[1,5-b]isoquinolinyl; or

_____ (7m) a C₆₋₁₄ aryl group optionally substituted by a C₁₋₆ alkoxy-
carbonyl group.

_____ 24. (New) The compound of claim 22, wherein R¹ and R² are the same
or different and each is a C₁₋₁₀ alkyl group optionally substituted by 1 to 3
substituent(s) selected from a C₃₋₁₀ cycloalkyl group, a C₁₋₆ alkoxy-carbonyl group
and a C₁₋₆ alkoxy group.

_____ 25. (New) The compound of claim 22, wherein R3 is a C6-14 aryl group optionally substituted by 1 to 3 substituent(s) selected from a C1-6 alkyl group optionally substituted by 1 to 3 halogen atom(s) and a halogen atom.

_____ 26. (New) The compound of claim 22, wherein Q is a bond.

_____ 27. (New) The compound of claim 1, wherein X is

_____ (3) (3a) a carboxyl group;

_____ (3b) a carbamoyl group;

_____ (3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy group;

_____ (3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3f) a C₇₋₁₃ aralkyloxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group, a halogen atom, a cyano group, a nitro group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom and a C₁₋₆ alkoxy group;

_____ (3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

_____ (3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

_____ (3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3m) a C₁₋₆ alkylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3n) a C₆₋₁₄ arylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkylsulfonyl group;

_____ (3o) a nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3p) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3q) a C₇₋₁₃ aralkyl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3r) a non-aromatic heterocycloxy-carbonyl group;

_____ (3s) a phosphono group optionally mono- or di-substituted by a C₁₋₆ alkyl group;

_____ (3t) an aromatic heterocyclyl-C₇₋₁₃ aralkyloxy-carbonyl group;

_____ (3u) a C₃₋₁₀ cycloalkyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (3v) a C₆₋₁₄ aryl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from an amino group optionally mono- or di-substituted by

a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group; or

_____ (3w) an aromatic heterocyclyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4) (4a) a C₁₋₆ alkyl-carbonyloxy group;

_____ (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (4d) a 5- or 6-membered aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4e) a fused aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4f) an aromatic heterocyclyl-C₁₋₆ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (4g) an aromatic heterocyclyl-C₆₋₁₄ aryloxy group;

_____ (5) (5a) a C₁₋₆ alkylthio group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (5b) a C₆₋₁₄ arylthio group optionally substituted by 1 to 3

substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group and a carbamoyl group; or

(5c) a 5- or 6-membered aromatic heterocyclylthio group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

(6) (6a) an amino group;

(6b) a C₁₋₆ alkoxy-carbonyl-C₁₋₁₀ alkylamino group;

(6c) a carboxy-C₁₋₁₀ alkylamino group;

(6d) a C₇₋₁₃ aralkyloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

(6e) a carbamoylamino group;

(6f) a mono- or di-C₁₋₆ alkyl-carbamoylamino group;

(6g) a C₁₋₆ alkylsulfonylamino group;

(6h) a C₆₋₁₄ arylsulfonylamino group optionally substituted by a C₁₋₆ alkylsulfonyl group;

(6i) an aromatic heterocyclyl-sulfonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group and a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

(6j) a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

(6k) a C₃₋₁₀ cycloalkyl-carbonylamino group;

(6l) a C₆₋₁₄ aryl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a cyano group, an optionally halogenated C₁₋₆ alkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group;

(6m) a C₇₋₁₃ aralkyl-carbonylamino group;

(6n) a C₈₋₁₃ arylalkenyl-carbonylamino group;

(6o) an aromatic heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a C₆₋₁₄ aryl group, a C₇₋₁₃ aralkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-

carbonyl group and a carbamoyl group;

_____ (6p) a nitrogen-containing heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6q) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonylamino group;

_____ (6r) a tetrahydropyranylcabonylamino group;

_____ (6s) a 4-oxo-4,5,6,7-tetrahydro-1-benzofuranyl-carbonylamino group;

_____ (6t) a C₁₋₆ alkoxy-carbonylamino group optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (6u) a C₆₋₁₄ aryloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6v) a C₇₋₁₃ aralkyl-carbamoylamino group; or

_____ (6w) an aromatic heterocyclyl-carbamoylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group.

28. (New) The compound of claim 22, wherein X is a carboxyl group.

29. (New) The compound of claim 22, which is 5-(aminomethyl)-2-methyl-4-(4-methylphenyl)-6-neopentyl nicotinic acid;
5-(aminomethyl)-6-isobutyl-2-methyl-4-(4-methylphenyl) nicotinic acid;
methyl 3-([5-(aminomethyl)-6-isobutyl-2-methyl-4-(4-methylphenyl)pyridin-3-yl]methoxy)-1-methyl-1H-pyrazole-4-carboxylate;
{[2-isobutyl-6-methyl-4-(4-methylphenyl)-5-(2-morpholin-4-yl-2-oxoethyl)pyridin-3-yl]methyl}amine;
methyl 3-([5-(aminomethyl)-6-isobutyl-2-methyl-4-(4-methylphenyl)pyridin-3-yl]acetyl)amino benzoate;

N-[5-(aminomethyl)-6-isobutyl-2-methyl-4-(4-methylphenyl)pyridin-3-yl]isoxazole-4-carboxamide,
or a salt thereof.

30. (New) A pharmaceutical agent comprising a compound of claim 22
or a salt thereof.

31. (New) The pharmaceutical agent of claim 30, which is an agent for
the prophylaxis or treatment of diabetes, diabetic complications, impaired
glucose tolerance or obesity.

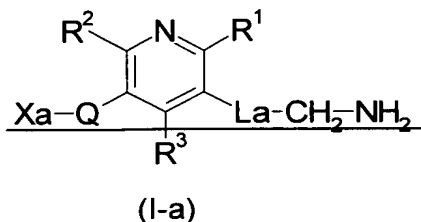
32. (New) A peptidase inhibitor comprising a compound of claim 22 or
a salt thereof.

33. (New) The inhibitor of claim 32, wherein the peptidase is dipeptidyl
dipeptidase-IV.

34. (New) A method for the prophylaxis or treatment of diabetes,
diabetic complications, impaired glucose tolerance or obesity in a mammal,
which comprises administering a compound of claim 22 or a salt thereof to the
mammal.

35. (New) A method of inhibiting peptidase in a mammal, which
comprises administering a compound of claim 22 or a salt thereof to the
mammal.

36. (New) A production method of a compound represented by the
formula



wherein

R¹, R², R³ and Q are as defined in claim 22;

La is a bond or a C₁₋₉ alkylene group; and

Xa is

_____ (3) (3a) a carboxyl group;

_____ (3b) a carbamoyl group;

_____ (3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3
substituent(s) selected from a carboxyl group, a carbamoyl group, a
thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy
group;

_____ (3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally
substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl
group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group
optionally substituted by a C₁₋₆ alkyl group;

_____ (3f) a C₇₋₁₃ aralkyloxy-carbonyl group optionally substituted by 1 to
3 substituent(s) selected from a carboxyl group, a carbamoyl group, a
thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group, a halogen atom, a cyano
group, a nitro group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group and a C₁₋₆
alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s)
selected from a halogen atom, a carboxyl group, C₁₋₆ alkoxy-carbonyl group and
a carbamoyl group);

_____ (3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl
group optionally substituted by 1 to 3 substituent(s) selected from a halogen
atom and a C₁₋₆ alkoxy group;

_____ (3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-
substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally
substituted by a C₁₋₆ alkyl group;

_____ (3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally
substituted by a C₁₋₆ alkyl group;

_____ (3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3

substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

_____ (3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3m) a C₁₋₆ alkylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3n) a C₆₋₁₄ arylsulfonyl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkylsulfonyl group;

_____ (3o) a nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (3p) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3q) a C₇₋₁₃ aralkyl-nitrogen-containing heterocyclyl-carbonyl group optionally substituted by 1 to 3 halogen atom(s);

_____ (3r) a non-aromatic heterocycloxy-carbonyl group;

_____ (3s) a phosphono group optionally mono- or di-substituted by a C₁₋₆ alkyl group;

_____ (3t) an aromatic heterocyclyl-C₇₋₁₃ aralkyloxy-carbonyl group;

_____ (3u) a C₃₋₁₀ cycloalkyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (3v) a C₆₋₁₄ aryl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from an amino group optionally mono- or di-substituted by a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group; or

_____ (3w) an aromatic heterocyclyl-carbamoyl group optionally

substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4) (4a) a C₁₋₆ alkyl-carbonyloxy group;

_____ (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

_____ (4d) a 5- or 6-membered aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4e) a fused aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (4f) an aromatic heterocyclyl-C₁₋₆ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (4g) an aromatic heterocyclyl-C₆₋₁₄ aryloxy group;

_____ (5) (5a) a C₁₋₆ alkylthio group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (5b) a C₆₋₁₄ arylthio group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group and a carbamoyl group; or

_____ (5c) a 5- or 6-membered aromatic heterocyclylthio group optionally

substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6) _____ (6a) an amino group;

_____ (6b) a C₁₋₆ alkoxy-carbonyl-C₁₋₁₀ alkylamino group;

_____ (6c) a carboxy-C₁₋₁₀ alkylamino group;

_____ (6d) a C₇₋₁₃ aralkyloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6e) a carbamoylamino group;

_____ (6f) a mono- or di-C₁₋₆ alkyl-carbamoylamino group;

_____ (6g) a C₁₋₆ alkylsulfonylamino group;

_____ (6h) a C₆₋₁₄ arylsulfonylamino group optionally substituted by a C₁₋₆ alkylsulfonyl group;

_____ (6i) an aromatic heterocyclyl-sulfonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group and a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

_____ (6j) a mono- or di-(C₁₋₆ alkyl-carbonyl)-amino group;

_____ (6k) a C₃₋₁₀ cycloalkyl-carbonylamino group;

_____ (6l) a C₆₋₁₄ aryl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a cyano group, an optionally halogenated C₁₋₆ alkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group;

_____ (6m) a C₇₋₁₃ aralkyl-carbonylamino group;

_____ (6n) a C₈₋₁₃ arylalkenyl-carbonylamino group;

_____ (6o) an aromatic heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a C₆₋₁₄ aryl group, a C₇₋₁₃ aralkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6p) a nitrogen-containing heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the

C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6q) a C₆₋₁₄ aryl-nitrogen-containing heterocyclyl-carbonylamino group;

_____ (6r) a tetrahydropyranylcabonylamino group;

_____ (6s) a 4-oxo-4,5,6,7-tetrahydro-1-benzofuranyl-carbonylamino group;

_____ (6t) a C₁₋₆ alkoxy-carbonylamino group optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (6u) a C₆₋₁₄ aryloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

_____ (6v) a C₇₋₁₃ aralkyl-carbamoylamino group; or

_____ (6w) an aromatic heterocyclyl-carbamoylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

_____ (7) (7a) tetrazolyl;

_____ (7b) oxoimidazolidinyl;

_____ (7c) dioxoimidazolidinyl optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7d) oxopiperazinyl;

_____ (7e) dioxopiperazinyl;

_____ (7f) oxodihydrooxadiazolyl;

_____ (7g) dioxoisindolyl;

_____ (7h) oxazolyl optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

_____ (7i) dioxooxazolidinyl or dioxothiazolidinyl, each of which is optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7j) 4-oxo-2-thioxo-1,3-thiazolidin-5-yl or 4-oxo-2-thioxo-1,3-

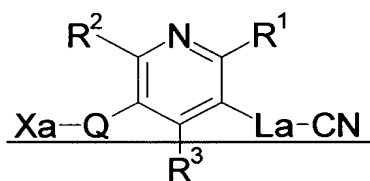
oxazolidin-5-yl, each of which is optionally substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group and a C₁₋₆ alkoxy-carbonyl group;

_____ (7k) 1,3(2H,5H)-dioxo-tetrahydroimidazo[1,5-a]pyridinyl;

_____ (7l) 1,3(2H,5H)-dioxo-10,10a-dihydroimidazo[1,5-b]isoquinolinyl; or

_____ (7m) a C₆₋₁₄ aryl group optionally substituted by a C₁₋₆ alkoxy-carbonyl group;

or a salt thereof, which comprises subjecting a compound represented by the formula



(II)

wherein each symbol is as defined above, or a salt thereof to a reduction reaction.

37. (New) The compound of claim 22, wherein R³ is a phenyl group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s) and a halogen atom.

38. (New) The compound of claim 22, wherein X is

(3) (3a) a carboxyl group;

(3b) a carbamoyl group;

(3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy group;

(3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by a C₁₋₆ alkyl group;

(3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom and a C₁₋₆ alkoxy group;

(3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

(3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

(3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

(3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

(3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(3r) a non-aromatic heterocyclyloxy-carbonyl group;

(4) (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group);

(4d) a 5- or 6-membered aromatic heterocyclyloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a

C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

(6) (6d) a C₇₋₁₃ aralkyloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

(6l) a C₆₋₁₄ aryl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a cyano group, an optionally halogenated C₁₋₆ alkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group;

(6o) an aromatic heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a C₆₋₁₄ aryl group, a C₇₋₁₃ aralkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

(6p) a nitrogen-containing heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group.

39. (New) The compound of claim 22, wherein X is

(3) (3a) a carboxyl group;

(3b) a carbamoyl group;

(3c) a C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl-carbonyloxy group;

(3d) an aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group, a thiocarbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by a C₁₋₆ alkyl group;

(3g) a carbamoyl group mono- or di-substituted by a C₁₋₆ alkyl

group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom and a C₁₋₆ alkoxy group;

(3h) a carbamoyl-C₁₋₆ alkyl-carbamoyl group optionally mono- or di-substituted by a C₁₋₆ alkyl group optionally substituted by 1 to 3 halogen atom(s);

(3i) a C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

(3j) a mono- or di-C₃₋₁₀ cycloalkyl-carbamoyl group optionally substituted by a C₁₋₆ alkyl group;

(3k) a C₇₋₁₃ aralkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a hydroxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a C₁₋₆ alkyl group;

(3l) an aromatic heterocyclyl-C₁₋₆ alkyl-carbamoyl group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group; or

(3r) a non-aromatic heterocyclyloxy-carbonyl group.

40. (New) The compound of claim 22, wherein X is

(3) (3a) a carboxyl group;

(3e) a non-aromatic heterocyclyl-C₁₋₆ alkoxy-carbonyl group optionally substituted by a C₁₋₆ alkyl group; or

(3r) a non-aromatic heterocyclyloxy-carbonyl group.

41. (New) The compound of claim 22, wherein X is

(4) (4b) a C₁₋₁₀ alkoxy group optionally substituted by 1 to 3 substituent(s) selected from a hydroxy group, a carboxyl group, a carbamoyl group and a C₁₋₆ alkoxy-carbonyl group;

(4c) a C₆₋₁₄ aryloxy group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, a C₁₋₆ alkylthio group, a carbamoyl group, a C₁₋₆ alkoxy group, a C₁₋₆ alkylsulfonyl group, a C₁₋₆ alkylsulfinyl group and a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group); or

(4d) a 5- or 6-membered aromatic heterocycloxy group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 or 2 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group.

42. (New) The compound of claim 22, wherein X is

(6) (6d) a C₇₋₁₃ aralkyloxy-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group;

(6l) a C₆₋₁₄ aryl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a halogen atom, a cyano group, an optionally halogenated C₁₋₆ alkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group, an aromatic heterocyclic group, a non-aromatic heterocyclic group and a carbamoyl group;

(6o) an aromatic heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group, a C₆₋₁₄ aryl group, a C₇₋₁₃ aralkyl group, a C₁₋₆ alkoxy group, a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group; or

(6p) a nitrogen-containing heterocyclyl-carbonylamino group optionally substituted by 1 to 3 substituent(s) selected from a C₁₋₆ alkyl group (the C₁₋₆ alkyl group is optionally substituted by 1 to 3 substituent(s) selected from a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group), a carboxyl group, a C₁₋₆ alkoxy-carbonyl group and a carbamoyl group.